

Report

Isocitrate dehydrogenase mutation is frequently observed in giant cell tumor of bone

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Key words

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Giant cell tumor of bone (GCTB) accounts for 5% of all primary bone tumors in adults in the USA and 20% in Asia.^(1,2) Although GCTB is generally benign, atypical GCTB may be associated with multiple local recurrences, multicentricity, and pulmonary metastases.⁽³⁾ Characterized by the presence of numerous multinucleated osteoclast-like giant cells, GCTB also includes mesenchymal fibroblast-like stromal cells and a mononuclear cell of myeloid lineage. Fibroblast-like stromal cells are considered to be responsible for the neoplastic character of the GCTB. The presence of telomeric associations, chromosomal aberrations, varied ploidy states, and gene amplifications have all been described within GCTB stromal cells.⁽⁴⁾ These stromal cells play an important role in the recruitment of tumor-associated myeloid lineage cells and formation of osteoclast-like giant cells.⁽⁵⁾ The stromal cells also induce osteoclastogenesis *in vitro* in coculture studies with osteoclasts, and produce several factors that are involved in the recruitment and induction of

Giant cell tumors of bone (GCTB) are benign and locally destructive tumors that include osteoclast-type multinuclear giant cells. No available treatment is definitively effective in curing GCTB, especially in surgically unresectable cases. Isocitrate dehydrogenase (IDH) mutations have been reported not only in gliomas and acute myeloid leukemias, but also in cartilaginous tumors and osteosarcomas. However, IDH mutations in GCTB have not been investigated. The IDH mutations are remarkably specific to arginine 132 (R132) in IDH1 and arginine 172 (R172) or arginine 140 (R140) in IDH2; IDH1/2 mutations are known to convert α -ketoglutarate to oncometabolite R(-)-2-hydroxyglutarate. We recently reported that the most frequent IDH mutation in osteosarcomas is IDH2-R172S, which was detected by MsMab-1, a multispecific anti-IDH1/2 mAb. Herein, we newly report the IDH mutations in GCTB, which were stained by MsMab-1 in immunohistochemistry. DNA direct sequencing and subcloning identified IDH mutations of GCTB as IDH2-R172S (16 of 20; 80%). This is the first report to describe IDH mutations in GCTB, and MsMab-1 can be anticipated for use in immunohistochemical determination of IDH1/2 mutation-bearing GCTB.

osteoclast differentiation and activation, including receptor activator of nuclear factor κ B ligand, the master regulator of osteoclast differentiation.⁽⁶⁾ Recently, it was reported that genes encoding histone H3.3 are frequently mutated in GCTB (92%).⁽⁷⁾

Isocitrate dehydrogenase (IDH) catalyzes the oxidative carboxylation of isocitrate to α -ketoglutarate.⁽⁸⁾ Mutated IDH1 and IDH2 convert α -ketoglutarate to oncometabolite R(-)-2-hydroxyglutarate (2-HG) in cytosol and mitochondria, respectively. Isocitrate dehydrogenase 1/2 mutations have been reported in gliomas,⁽⁹⁾ acute myeloid leukemias,⁽¹⁰⁾ cartilaginous tumors,⁽¹¹⁾ osteosarcomas,⁽¹²⁾ Ollier disease,⁽¹¹⁾ and Maffucci syndrome.^(11,13) The heterozygous IDH mutations are remarkably specific to a single codon in the conserved and functionally important arginine 132 residue (R132) of IDH1 and 172 residue (R172) of IDH2. We have established multi-specific anti-IDH1/2 mAbs^(14,15) that are useful for diagnosis of IDH1/2 mutation-bearing tumors. Herein, we report the

Table 1. The characteristic of giant cell tumor patients used in immunohistochemical analysis by MsMab-1

Patient no.	Age	Gender	Race	Sample class	Site	MsMab-1 staining (Mesenchymal stromal cells)		IDH1 (R132)	IDH2 (R172)		IDH2 (H175)	H3F3A (K27, G34, K36)	H3F3B (G34, K36)	
						Percentage†	Intensity‡		Direct sequencing					Subcloning
1	32	M	Asian	Primary	Tibia	+++	+	WT	R172S	WT	WT	WT		
2	17	F	Asian	Primary	Femur	+	+	WT	R172S	WT	WT	WT		
3	42	F	Asian	Primary	Humerus	±	+	WT	WT	R172S (2/25: 8%)	WT	WT		
4	60	F	Asian	Primary	Maxilla	±	+	WT	WT	R172S (1/7: 14%)	WT	WT		
5	24	F	Asian	Primary	Humerus	+++	+	WT	R172S	WT	WT	WT		
6	38	F	Asian	Primary	Radius	++	+	WT	WT	R172S (6/21: 29%)	WT	WT		
7	34	M	Asian	Primary	Tibia	+++	++	WT	R172S	WT	WT	WT		
8	45	F	Asian	Primary	Tibia	+++	++	WT	R172S	WT	WT	WT		
9	33	F	Asian	Primary	Femur	+++	+	WT	R172S	H175Y	WT	WT		
10	40	F	Asian	Primary	Radius	++	+	WT	R172S	WT	WT	WT		
11	33	M	Asian	Primary	Humerus	-	-	WT	WT	WT	WT	WT		
12	36	F	Asian	Primary	Tibia	+++	+	WT	R172S	WT	WT	WT		
13	28	M	Asian	Primary	Clavicle	++	+	WT	R172S	H175Y	WT	WT		
14	48	F	Asian	Primary	Femur	++	+	WT	R172S	WT	WT	WT		
15	23	M	Asian	Primary	Femur	±	+	WT	WT	R172S (0/38: 0%)	WT	WT		
16	34	M	Asian	Primary	Sacrum	±	+	WT	WT	R172S (0/42: 0%)	WT	WT		
17	50	M	Asian	Primary	Femur	++	+	WT	R172S	WT	WT	WT		
18	38	F	Asian	Primary	Humerus	±	+	WT	WT	R172S (0/41: 0%)	H175Y	WT		
19	47	M	Asian	Primary	Tibia	+	+	WT	R172S	H175Y	WT	WT		
20	20	M	Asian	Primary	Femur	++	+	WT	R172S	WT	WT	WT		

†-, no staining; ±, <1%; +, 1-10%; ++, 10-50%; and +++>50%. ‡-, no staining; +, weak; ++, medium; +++, strong.

IDH2-R172S mutation in GCTB patients, which was detected by MsMab-1 mAb and direct DNA sequencing.

Materials and Methods

Immunohistochemical analyses. Tissue microarrays (BO2081; US Biomax, Rockville, MD, USA) were used in this study. Immunohistochemical analyses were carried out as described in Document S1.

Direct DNA sequencing of IDH1, IDH2, H3F3A, and H3F3B. Genomic DNA extraction and PCR were carried out as described in Document S1.

Plasmid preparation, protein expression, and Western blot analyses. Osteosarcoma U-2 OS cells were transfected with appropriate amounts of plasmids as described in Document S1. The SDS-PAGE and Western blot analyses using MsMab-1 or anti-PA tag (NZ-1)^(14–16) were carried out as described in Document S1.

Analysis of 2-HG production. Sample preparation and measurement by capillary electrophoresis time-of-flight mass spectrometry are described in Document S1.

Results

Immunohistochemical analysis by MsMab-1 against GCTB. We carried out immunohistochemistry against GCTB using a multispecific antimutated IDH1/2 mAb, MsMab-1. The characteristics of the GCTB patients are presented in Table 1. Typical staining patterns are shown in Figure 1. Both multinucleated osteoclast-like giant cells and mesenchymal fibroblast-like stromal cells were diffusely stained by MsMab-1 (Fig. 1a,b). In contrast, weak and focal staining of mesenchymal fibroblast-

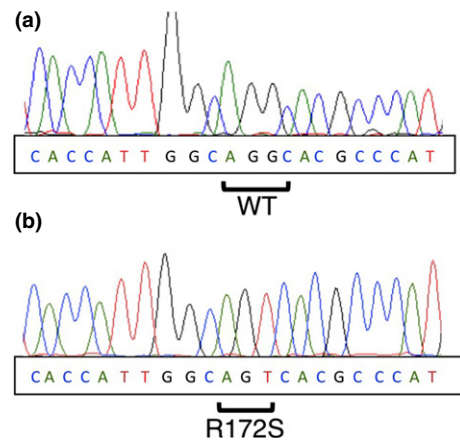


Fig. 2. Mutational analysis of isocitrate dehydrogenase 1/2 in giant cell tumor of bone. (a) DNA direct sequencing of giant cell tumor of sample no. 3. (b) Subcloning of PCR products.

like stromal cells was observed in other samples (Table 1). Because MsMab-1 stained multinucleated giant cells in foreign-body granulomas (Fig. S1), multinucleated osteoclast-like giant cells in GCTB might be non-specifically stained by MsMab-1 (Fig. 1).

Mutational analyses in GCTB. Polymerase chain reaction was carried out using DNA samples obtained from tissue microarray. No IDH1 mutation was observed in 20 samples (Table 1). In contrast, 13 of 20 (65%) GCTB samples possessed IDH2 mutations. It is noteworthy that all 13 IDH2 mutations were of IDH2-R172S (AGG > AGT; Fig. 1d,e), which is also frequently observed in osteosarcomas and

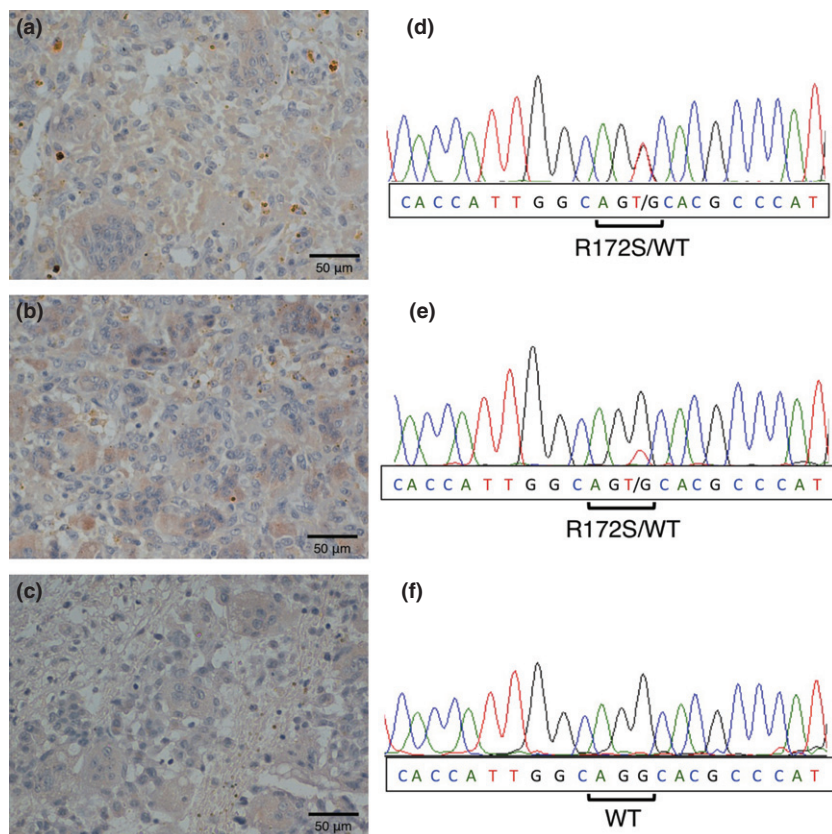


Fig. 1. Mutational analysis of isocitrate dehydrogenase 1/2 (IDH1/2) in giant cell tumor of bone. (a–c) Immunohistochemical analysis by MsMab-1, a multispecific anti-IDH1/2 mAb, against tissue microarray of giant cell tumor of bone. (d–f) DNA direct sequencing. (a, d) Sample no. 7; (b, e) no. 8; (c, f) no. 11.

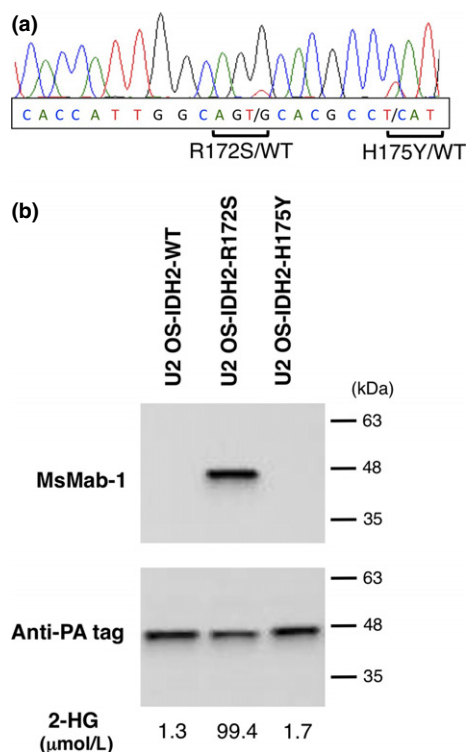


Fig. 3. Isocitrate dehydrogenase 2 (IDH2)-H175Y (CAT > TAT) mutations were not detected by MsMab-1, a multispecific anti-IDH1/2 mAb. (a) DNA direct sequencing was carried out against bone sample no. 19. (b) Total cell lysate from U-2 OS osteosarcoma cells expressing IDH2 wild-type-PA tag (WT, lane 1) and IDH2 mutants (lane 2, IDH2-R172S-PA tag; lane 3, IDH2-H175Y-PA tag) were electrophoresed, and Western blotted using MsMab-1 or anti-PA tag (NZ-1). Production of oncometabolite R(-)-2-hydroxyglutarate (2-HG) was observed in IDH2-R172S, but not in IDH2-H175Y or IDH2-WT.

chondrosarcomas.^(11,12) After subcloning of PCR products, 3 of 6 (50%) GCTB samples were shown to possess IDH2-R172S (Fig. 2, Table 1). In total, 16 of 20 (80%) GCTB samples were shown to possess IDH2-R172S (Table 1). In 5 of 20 (25%) GCTB patients, IDH2-H175Y (CAT > TAT) mutations were detected (Fig. 3a, Table 1), although IDH2-H175Y mutation was not recognized by MsMab-1 in Western blot analyses (Fig. 3b). The U2 OS IDH2-R172S cells produced 99.4 μmol/L of oncometabolite 2-HG, whereas U2 OS IDH2-H175Y, U2 OS IDH2-WT, and U2 OS cells produced 1.7, 1.3, and 1.6 μmol/L 2-HG, respectively (Fig. 3b).

Discussion

We recently reported that IDH mutations are observed in osteosarcomas.⁽¹²⁾ Herein, we investigated the IDH mutations in

References

- Turcotte RE. Giant cell tumor of bone. *Orthop Clin North Am* 2006; **37**: 35–51.
- Szendroi M. Giant-cell tumour of bone. *J Bone Joint Surg Br* 2004; **86**: 5–12.
- Balke M, Hards J. Denosumab: a breakthrough in treatment of giant-cell tumour of bone? *Lancet Oncol* 2010; **11**: 218–9.
- Schwartz HS, Eskew JD, Butler MG. Clonality studies in giant cell tumor of bone. *J Orthop Res* 2002; **20**: 387–90.

GCTB, because GCTB accounts for 5–20% of all primary bone tumors in adults.^(1,2) Although both multinucleated osteoclast-like giant cells and mesenchymal fibroblast-like stromal cells were stained by MsMab-1 (Fig. 1), multinucleated osteoclast-like giant cells might be non-specifically stained by MsMab-1, because MsMab-1 also stained giant cells in foreign-body granulomas (Fig. S1). MsMab-1 can recognize over-expressed wild-type IDH1/2 in Western blot analyses;⁽¹²⁾ therefore, osteoclast-like giant cells in GCTB might overexpress wild-type IDH1/2 proteins. To clarify this issue, we should develop an anti-IDH2-R172S-specific mAb in the near future. We analyzed IDH mutations using 20 GCTB specimens with direct DNA sequencing (Table 1). Thirteen of 20 (65%) GCTB samples possessed IDH2 mutations. Furthermore, PCR products of sample numbers 3, 4, and 6, included the IDH2-R172S mutation after subcloning, indicating that MsMab-1 indeed detected IDH2-R172S in those tissues (Table 1).⁽¹²⁾ The PCR products of sample numbers 15, 16, and 18 were not shown to possess IDH2-R172S mutation after subcloning; therefore, MsMab-1 reaction against these GCTB tissues might be non-specific, or little fraction of IDH2-R172S was included in these GCTB tissues. We will carefully check the MsMab-1 reaction in future immunohistochemical studies. Because the IDH2-H175Y mutation was not recognized by MsMab-1 in Western blot analyses (Fig. 3b), IDH2-H175Y is not relevant with MsMab-1 staining in immunohistochemistry. Furthermore, IDH2-H175Y did not produce oncometabolite 2-HG (Fig. 3b). We did not observe any clinical difference between IDH2 mutation-positive patients and IDH2 mutation-negative patients in this study; the number of patients should be increased to investigate the clinical importance of IDH2 mutation in GCTB in the future. We also investigated H3F3A and H3F3B mutations in the GCTB samples. However, we observed neither H3F3A mutations (K27, G34, K36) nor H3F3B mutations (G34, K36) in this study (Table 1, Figs S2, S3). We need further investigations to clarify the difference between this study and the previous one.⁽⁷⁾ Furthermore, anti-mutated H3F3A/H3F3B-specific mAbs could be useful for investigating H3F3A and H3F3B mutations in combination with antimutated IDH1/2 mAbs.^(14,15,17–23)

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Disclosure Statement

The authors have no conflicts of interest.

- Cowan RW, Singh G. Giant cell tumor of bone: a basic science perspective. *Bone* 2013; **52**: 238–46.
- Atkins GJ, Kostakis P, Vincent C et al. RANK expression as a cell surface marker of human osteoclast precursors in peripheral blood, bone marrow, and giant cell tumors of bone. *J Bone Miner Res* 2006; **21**: 1339–49.
- Behjati S, Tarpey PS, Presneau N et al. Distinct H3F3A and H3F3B driver mutations define chondroblastoma and giant cell tumor of bone. *Nat Genet* 2013; **45**: 1479–82.
- Dang L, White DW, Gross S et al. Cancer-associated IDH1 mutations produce 2-hydroxyglutarate. *Nature* 2009; **462**: 739–44.

- 9 Yan H, Parsons DW, Jin G *et al*. IDH1 and IDH2 mutations in gliomas. *N Engl J Med* 2009; **360**: 765–73.
- 10 Ward PS, Patel J, Wise DR *et al*. The common feature of leukemia-associated IDH1 and IDH2 mutations is a neomorphic enzyme activity converting alpha-ketoglutarate to 2-hydroxyglutarate. *Cancer Cell* 2010; **17**: 225–34.
- 11 Pansuriya TC, van Eijk R, d'Adamo P *et al*. Somatic mosaic IDH1 and IDH2 mutations are associated with enchondroma and spindle cell hemangioma in Ollier disease and Maffucci syndrome. *Nat Genet* 2011; **43**: 1256–61.
- 12 Liu X, Kato Y, Kaneko MK *et al*. Isocitrate dehydrogenase 2 mutation is a frequent event in osteosarcoma detected by a multi-specific monoclonal antibody MsMab-1. *Cancer Med* 2013; **2**: 803–14.
- 13 Moriya K, Kaneko MK, Liu X *et al*. IDH2 and TP53 mutations are correlated with gliomagenesis in a patient with Maffucci syndrome. *Cancer Sci* 2014; **105**: 359–62.
- 14 Kaneko M, Ogasawara S, Kato Y. Establishment of a novel multi-specific monoclonal antibody MsMab-1 recognizing both IDH1 and IDH2 mutations. *Tohoku J Exp Med* 2013; **230**: 103–9.
- 15 Ogasawara S, Kaneko MK, Tsujimoto Y, Liu X, Kato Y. A multi-specific monoclonal antibody MsMab-2 recognizes IDH1-R132L and IDH2-R172M mutations. *Monoclon Antib Immunodiagn Immunother* 2013; **32**: 377–81.
- 16 Fujii Y, Kaneko M, Neyazaki M, Nogi T, Kato Y, Takagi J. PA tag: a versatile protein tagging system using a super high affinity antibody against a dodecapeptide derived from human podoplanin. *Protein Expr Purif* 2014; **95**: 240–7.
- 17 Kato Y, Jin G, Kuan CT, McLendon RE, Yan H, Bigner DD. A monoclonal antibody IMab-1 specifically recognizes IDH1R132H, the most common glioma-derived mutation. *Biochem Biophys Res Commun* 2009; **390**: 547–51.
- 18 Kaneko MK, Tian W, Takano S, *et al*. Establishment of a novel monoclonal antibody SMab-1 specific for IDH1-R132S mutation. *Biochem Biophys Res Commun* 2011; **406**: 608–13.
- 19 Kaneko MK, Tsujimoto Y, Hozumi Y, Goto K, Kato Y. Novel monoclonal antibodies GMab-r1 and LMab-1 specifically recognize IDH1-R132G and IDH1-R132L mutations. *Monoclon Antib Immunodiagn Immunother* 2013; **32**: 224–8.
- 20 Kaneko MK, Morita S, Tsujimoto Y, *et al*. Establishment of novel monoclonal antibodies KMab-1 and MMab-1 specific for IDH2 mutations. *Biochem Biophys Res Commun* 2013; **432**: 40–5.
- 21 Kato Y, Kaneko MK. Generation of a novel monoclonal antibody WMab-1 specific for IDH2-R172W mutation. *Biochem Biophys Res Commun* 2013; **433**: 374–8.
- 22 Kato Y, Natsume A, Kaneko MK. A novel monoclonal antibody GMab-m1 specifically recognizes IDH1-R132G mutation. *Biochem Biophys Res Commun* 2013; **432**: 564–7.
- 23 Takano S, Kato Y, Yamamoto T, *et al*. Immunohistochemical detection of IDH1 mutation, p53, and internexin as prognostic factors of glial tumors. *J Neuro oncol* 2012; **108**: 361–73.

Supporting Information

Additional supporting information may be found in the online version of this article:

Fig. S1. Immunohistochemical analysis by MsMab-1 against inflammatory tissues.

Fig. S2. Mutational analysis of H3F3A in giant cell tumor of bone.

Fig. S3. Mutational analysis of H3F3B in giant cell tumor of bone.

Data S1. Detailed materials and methods.